

Resource Summary Report

Generated by [nidm-terms](#) on Aug 10, 2026

pJ4M/TDP-43

RRID:Addgene_104480

Type: Plasmid

Proper Citation

RRID:Addgene_104480

Plasmid Information

URL: <http://www.addgene.org/104480>

Proper Citation: RRID:Addgene_104480

Insert Name: TDP-43

Organism: Homo sapiens

Bacterial Resistance: Kanamycin

Defining Citation: [PMID:29438978](#)

Vector Backbone Description: Vector Backbone:pJ4M; Vector Types:Bacterial Expression; Bacterial Resistance:Kanamycin

Plasmid Name: pJ4M/TDP-43

Record Creation Time: 20220422T221507+0000

Record Last Update: 20260808T080333+0000

Ratings and Alerts

No rating or validation information has been found for pJ4M/TDP-43.

No alerts have been found for pJ4M/TDP-43.

Data and Source Information

Source: [Addgene](#)

Usage and Citation Metrics

We found 24 mentions in open access literature.

Listed below are recent publications. The full list is available at [nidm-terms](#) .

Guo L, et al. (2026) Defining RNA oligonucleotides that reverse deleterious phase transitions of RNA-binding proteins with prion-like domains. *Molecular cell*, 86(1), 114.

Copley KE, et al. (2026) Short RNA chaperones promote aggregation-resistant TDP-43 conformers to mitigate neurodegeneration. *Science (New York, N.Y.)*, 392(6798), eadv3301.

Copley KE, et al. (2026) Isoform-specific steric zippers drive aberrant assembly and mislocalization of shortened TDP-43. *Science advances*, 12(26), eady0256.

Elmaleh B, et al. (2026) The ALS-associated E425K mutation uncouples DNAJC7 from the Hsp70 chaperone cycle. *The FEBS journal*, 293(12), 3485.

Aikio M, et al. (2025) Opposing roles of p38 γ -mediated phosphorylation and PRMT1-mediated arginine methylation in driving TDP-43 proteinopathy. *Cell reports*, 44(1), 115205.

Oh J, et al. (2025) Engineering a membrane protein chaperone to ameliorate the proteotoxicity of mutant huntingtin. *Nature communications*, 16(1), 737.

Zibold J, et al. (2024) The new missense G376V-TDP-43 variant induces late-onset distal myopathy but not amyotrophic lateral sclerosis. *Brain : a journal of neurology*, 147(5), 1768.

Hayashi M, et al. (2024) Engineered NLS-chimera downregulates expression of aggregation-prone endogenous FUS. *Nature communications*, 15(1), 7887.

Minshull TC, et al. (2024) Hydrogen-Deuterium Exchange Mass Spectrometry Reveals Mechanistic Insights into RNA Oligonucleotide-Mediated Inhibition of TDP-43 Aggregation. *Journal of the American Chemical Society*, 146(49), 33626.

Lynch EM, et al. (2024) Seeding competent TDP-43 persists in human patient and mouse muscle. *bioRxiv : the preprint server for biology*.

Ervilha Pereira P, et al. (2023) C-terminal frameshift variant of TDP-43 with pronounced aggregation-propensity causes rimmed vacuole myopathy but not ALS/FTD. *Acta neuropathologica*, 145(6), 793.

Pérez-Berlanga M, et al. (2023) Loss of TDP-43 oligomerization or RNA binding elicits distinct aggregation patterns. *The EMBO journal*, 42(17), e111719.

Abayev-Avraham M, et al. (2023) DNAJB6 mutants display toxic gain of function through unregulated interaction with Hsp70 chaperones. *Nature communications*, 14(1), 7066.

Riemenschneider H, et al. (2023) Targeting the glycine-rich domain of TDP-43 with antibodies prevents its aggregation in vitro and reduces neurofilament levels in vivo. *Acta neuropathologica communications*, 11(1), 112.

Ma XR, et al. (2022) TDP-43 represses cryptic exon inclusion in the FTD-ALS gene UNC13A. *Nature*, 603(7899), 124.

Shen H, et al. (2022) Sexually dimorphic RNA helicases DDX3X and DDX3Y differentially regulate RNA metabolism through phase separation. *Molecular cell*, 82(14), 2588.

Peinado JR, et al. (2022) Sequestration of TDP-43²¹⁶⁻⁴¹⁴ Aggregates by Cytoplasmic Expression of the proSAAS Chaperone. *ACS chemical neuroscience*, 13(11), 1651.

Hallegger M, et al. (2021) TDP-43 condensation properties specify its RNA-binding and regulatory repertoire. *Cell*, 184(18), 4680.

Perdikari TM, et al. (2020) SARS-CoV-2 nucleocapsid protein undergoes liquid-liquid phase separation stimulated by RNA and partitions into phases of human ribonucleoproteins. *bioRxiv : the preprint server for biology*.

Perdikari TM, et al. (2020) SARS-CoV-2 nucleocapsid protein phase-separates with RNA and with human hnRNPs. *The EMBO journal*, 39(24), e106478.